

# **Product Data Sheet**

### Anti-14-3-3 beta Antibody

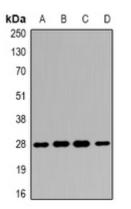
Catalog #	Source	Reactivity	Applications
CQA2325	Rabbit	H, M, R	WB, IH, IF/IC
Description	R	abbit polyclonal antibody	to 14-3-3 beta
Immunogen	R	Recombinant full length pro	tein of human 14-3-3 beta
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.
Specificity	R	Recognizes endogenous lev	els of 14-3-3 beta protein.
Clonality	Р	Polyclonal	
Conjugation			
Form	L	iquid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	а	nd 0.01% sodium azide.	
Dilution	V	VB (1/500 - 1/2000), IH (1/50	) - 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	Y	(WHAB	
Alternative N	ames 1	.4-3-3 protein beta/alpha;	Protein 1054; Protein kinase C inhibitor protein 1; KCIP-1
Entrez Gene	7	2529 (Human); 54401 (Μοι	se); 56011 (Rat)
SwissProt	Р	231946 (Human); Q9CQV8	(Mouse); P35213 (Rat)
Storage/Stabi	i <b>lity</b> S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	fi	reeze/thaw cycles.	

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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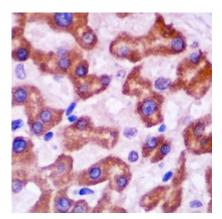




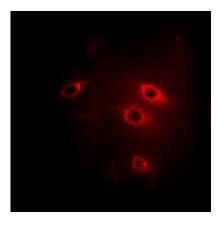
For research purposes only, not for human use

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Western blot analysis of 14-3-3 beta expression in SW480 (A), HepG2 (B), mouse liver (C), mouse kidney (D) whole cell lysates. (Predicted band size: 27; 28 kD; Observed band size: 28 kD)



Immunohistochemical analysis of 14-3-3 beta staining in rat liver formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of 14-3-3 beta staining in Hela cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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